HFS Prior Approval Information Form for Biologic Modifier agents http://www.hfs.illinois.gov/pharmacy/guidelines.html

A. PHYSICIAN INFORMATION ALL Information Requested On This Form Must Be Complete					
Physician Name:	DEA #:			License #:	
	NO				
Prescriber is a Rheumatologist?	(If NO, list specialty)			Office phone #:	
B. PATIENT INFORMATION					
Patient Name:	DOB//	Patient 9 d	ligit IDPA	Recipient Number:	
			Ü	•	
C. PATIENT INFORMATION - DIAGNOSIS	and THERAPY INFORMATION				
Diagnosis:	Current Patient W	eight:			
Please complete the following EVEN V	VITH RENEWAL REQUEST	YES	NO	Ι	
Failure of at least two forms of DMARD therapy (if	no, explain in comment section)			REQUESTED DOSE/NDC	
Positive PPD prior to initiating biologic agent therapy					
History or pre-existing neurological diseases (incl. demyelinating disorders)				☐ Enbrel 25mg: 58406042534 twice weekly	
Lupus erythematosus or lupus-like syndrome			☐ Enbrel 50mg: 58406043501 twice weekly		
History of malignancy			Emplei Soing. 30400043301 twice weekly		
Evidence or history of heart failure			Humira 40mg: 00074379902 every other wk		
Evidence of infection			Itumina 40mg. 000/43/7702 every other wk		
Patient is educated on the administration of the med			Remicade 100mg: 57894003001 Dose =		
Associated risks and monitoring for side effects (e.g			Orencia 250 mg: 00003218710 Dose =		
Is this a change in agent or dose? If yes, explain in S			Grencia 230 mg. 00003210710 Bosc =		
D. DRUG ACQUISITION and ADMINISTRA	TION INFORMATION				
		YES	NO	Pharmacy name:	
Medication will be dispensed to patient by retail pha	ırmacy			I narmacy name.	
Medication will be dispensed to Long-Term Care (LTC) facility by LTC pharmacy]., , , ,	
				Pharmacy phone #:	
E. ADDITIONAL INFORMATION					
IMPORTANT: To prevent delay, fax relevant	patient information with this form t	o validate reg	uest, or li	st comments below	
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FAX TO: 217-524-7264 ATTN: MEDICAL COMMITTEE

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F. PHYSICIAN or DESIGNEE'S SIGNATURE:	Date:
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